PATENT SPECIFICATION

NO DRAWINGS



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We, Dow Corning Corporation, of Midland, Michigan, United States of America, a corporation organised under the laws of the State of Michigan, United States 5 of America, do hereby declare the invention, for which we pray that a patent may be granted to us, and the method by which it is to be performed, to be particularly described in and by the following statement:-

This invention relates to a method of imparting durable creases and crease resistance

to garments containing thermoplastic fibres. It is known that so-called permanent creases (i.e., creases which are resistant to launder-15 ing or dry cleaning) can be imparted to cotton fabric by impregnating the fabric with an aqueous solution of an urea- or melamineformaldehyde resin, drying the fabric, forming it into a garment and then pressing the 20 garment to form creases in the desired loca-

The pressing is carried out under conditions of heat and pressure which cause the formaldehyde resin to cure while the garment is held in the creased position, thereby fixing the crease in such a way that it is not obliterated by washing or normal wear.

It is also known that silicones can be used in conjunction with these formaldehyde resins 30 in order to soften the finished fabric, thereby imparting a better hand. The silicone is used as a coating on the fabric which has been impregnated with the permanent crease forming resin.

It has unexpectedly been found according to the invention the silicones alone will impart a durable crease to certain types of fabrics, as shown below.

It is an object of this invention to provide permanent creases in garments by a method which obviates the use of heretofore employed permanent crease-forming resins so that one and water-repellency in one operation and with only one treatement.

This invention provides a process of imparting permanent creases to a garment which comprises:

(1) applying to a fabric comprising fibres formed from a thermoplastic organic polymer a curable organosiloxane in an amount of at least 0.5% by weight calculated on the weight of the fabric,

(2) forming the fabric into a garment, and (3) arranging those areas of the garment in which it is desired to form creases in creases and subjecting the garment to sufficient heat and pressure whilst so arranged so as to set the thermoplastic fibres and cure the siloxane to a flexible resin, said siloxane being the only permanent crease-forming resin applied to the fabric.

The term "permanent crease-forming resin" as employed herein means a resinous composition which is applied to the fabric for the purpose of imparting a permanent crease to the fabric.

For the purposes of this invention the fabric can be made from any thermoplastic organic fibre. As is well-known, these fibres are generally melt-spun and include, for example, fibres made of synthetic plastics, such as polyesters, such as polyesters including long chain synthetic polymers composed of an ester of teraphthalic acid and ethylene glycol; polyamides such as long chain synthetic polyamides having recurring amide groups as an integral part of the polymer chain, e.g. nylon polyacrylonitrile fibres; co-polymers of vinyl chloride and vinylidene chloride; poly-vinylidenenitrile; polyethylene; polypropylene; copolymers of any of the above; ethyl cellulose and cellulose acetate. The term "fabric comprising fibres formed from a

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[Price 4s. 6d.]

thermoplastic organic polymer" also includes blends of those thermoplastic fibres with natural fibres such as cotton, wool, silk and linen, in which the thermoplastic fibre makes up a major portion (i.e. at least 35% and preferably at least 50%) of the fibres in the

The silicones applicable to this invention are any of those which cure to flexible As is well-known these, in materials. general, comprise the di-substituted organopolysiloxanes in which the substituents are essentially all hydrocarbon radicals or halogenated hydrocarbon radicals such as aliphatic hydrocarbon radicals such as ethyl, methyl, propyl, octadecyl, vinyl, allyl and hexenyl; cycloaliphatic hydrocarbon radicals such as cyclohexyl, cyclopentyl, cyclohexenyl and methylcyclopentyl; aromatic hydrocarbon radicals such as phenyl, xenyl, tolyl, and naphthyi, and aralkyl hydrocarbon radicals such as benzyl, β -phenylethyl and β -phenylpropyl. The substituents can also be any halogenated hydrocarbon radical such as chloromethyl, gammachloropropyl, chlorophenyl, bromophenyl, chloroxenyl, bromocyclohexyl, a,a,a,-trifluorotolyl and radicals of the general formula RCH2CH2-, in which R is a perfluoroalkyl radical such as trifluoromethyl, pentafluoroethyl and C,F15.

The siloxane can be cured on the fabric with any of the conventional curing techniques employed by siloxanes. For example, the siloxanes can be cured with peroxide catalysts or any of the standard catalysts normally employed with siloxane materials. Often it is desirable to employ siloxane crosslinking materials which aid in the cure of the siloxane and these include methyl hydrogen siloxane, alkyl silicates such as ethylsilicate; acyloxy silanes such as methyltriacetoxy silane C_2H_3Si oxime silanes such as

 $[ON = C(CH_3)_2]_3.$ After the siloxane has been appled to the fabric and the fabric formed into a garment, those areas of the garment in which it is desired to form creases are arranged in creases and the garment subjected to heat and pressure. The amount of heat needed should be sufficient to set the thermo plastic fibre and to cure the siloxane resin. In general, this can be accomplished at the temperature of 150° to 205°C., although these values are not critical and will vary, depending on the nature of the thermoplastic fibres or the proportion of the thermoplastic fibre in the fabric and on the nature of the siloxane. The amount of pressure need be only that sufficient to form a satisfactory crease, and in 60 general, 700 to 1050 g./sq cm. is sufficient.

The garment can be held in the press for a sufficient time to cure the siloxane completely. However, this is a rather slow process and it is generally desirable for economic reasons to hold the garment in the press for

sufficient time to form the crease and partially to cure the siloxane, thereby setting the crease and then to place the garment in a heated environment, such as an oven, in order to complete the cure of the siloxane. This enables a faster operation of the pross.

With most siloxanes a press-cure of 15 seconds at 150° to 205°C. and 700 to 1050 g./sq cm. of pressure and a post-cure of some 4 to 15 minutes at 150° to 205°C. is sufficient. It should be understood, of course, that these values are not critical.

If desired, other materials can be incorporated in the siloxane resin in order to perform specific functions not related to crease resistance. For example, small amounts of anchoring agents such as vinylacetoxysilanes can be included in order to increase the adhesion of the silexane to the fabric. In addition one can incorporate other materials in small amounts to reduce undesirable odours which may arise, or one can incorporate sun screen agents or other materials for incidental reasons not related to the crease phenomenon.

The siloxane can be applied to the fabric in any desirable manner such as from a solution or an emulsion by dipping, spraying or padding. It is necessary to apply at least 0.5% by weight of siloxane solids to the fabric in order to obtain the desired crease performance. The upper limit of siloxane is not critical. For economic reasons one would not normally employ more than 5% by weight.

The following Examples illustrate the in- 100 vention. All percents are by weight.

Example 1

A fabric composed by 65% of a polyester fibre comprising a co-polymer of terephthalic acid and ethylene glycol and 35% of cotton 105 was padded with a solution having the following composition

4.3% of a polydimethylsiloxane gum containing 3% by weight of methyl hydrogen polysiloxane,

0.16% of dibutyl tin dilaurate 0.5% of vinyltriacetoxysilane

0.16% of additional methyl hydrogen polysiloxane

0.05% of a trimethyl-end-blocked poly- 115 dimethyl siloxane of 60,000cs. viscosity at 25°C.,

25°C., 0.1% of ethylpolysilicate

0.2% of an urea-formaldehyde odour reduc- 120 ing agent

0.1% of acetic acid and 94.43% of perchloroethylene.

The fabric was dried to remove the solvent and the silicone pickup was 1.5% by weight. 125 The fabric was sewn into trousers, and the

trousers were pressed for 15 seconds at 171°C. at a pressure of 700 to 1050 g./sq.cm. steam in order to form a crease in the

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1,123,447 desired position. The trousers were then 0.26% of a mixture of dibutyl tin diacetate 60 removed from the press and heated for 10 and zinc octoate, the remainder being to 15 minutes at 163°C. The resulting garment was washed five The silicone pick-up was 1.5 per cent by times at 60°C. in a conventional household washer employing a commercial detergent. The fabric was dried, sewn into trousers and The trousers retained their crease and the pressed as shown in Example 2, and then heated for 10 minutes at 163°C. The fabric fabric had an excellent hand. showed excellent crease retention after being Example 2 washed five times as shown in Example 2. A fabric as defined in Example 1 was padded with the following composition: Example 5 70 1.56% of a mixture of 35.9 parts by weight of Excellent crease retention was obtained a dimethyl-polysiloxane gum, 23.2 parts when fabrics made from the following types by weight of a hydroxylated dimethylpolyof fibres were substituted in the procedure of siloxane liquid and 1.8 parts by weight of Example 2; co-polymers of vinyl chloride and methyl polysiloxane, vinylidene chloride, polyethylene, polypropy-0.3% of dibutyl tin dilaurate lene, cellulose acetate and polyacrylonitrile. 0.5% of vinyltriacetoxysilane In each case the silicone pick-up was 1.5 0.1% of acetic acid per cent by weight. 0.5% of tetrabutyltitanate the remainder being perchloroethylene. Example 6 Improved crease retention was obtained The silicone pick-up was 1.3 per cent by when the following siloxane gums were subweight. stituted for the dimethylpolysiloxane gum of The fabric was dried and allowed to stand Example 2: for 20 days at the end of which time it was sewn in to trousers and was pressed under the 1. a co-polymer of dimethylsiloxane and conditions shown in Example 1, and therephenylmethylsiloxane 85 after removed from the press and heated for a co-polymer of dimethylsiloxane and di-10 minutes at 163°C. The fabric showed phenylsiloxane excellent crease retention after five washings 3. polyethylmethylsiloxane at 60°C. with a commercial laundry detergent 4. polymethylvinyl siloxane in a household washing machine with tumble 5. a co-polymer of dimethylsiloxane and drying. octadecylmethylsiloxane 6. a co-polymer of dimethylsiloxane and Example 3 chlorophenyl-methyl-siloxane. The fabric of Example 2 was padded with 7. a co-polymer of β -phenylpropyl(methyl)the following composition: siloxane and dimethylsiloxane and 8. poly (3,3,3-trifluoropropyl)methylsiloxane. 3% of a mixture of 97% of a dimethylpoly-In each case the silicone pick-up was 1.5 silexane gum and per cent by weight. 3% of methyl hydrogen polysiloxane 0.5% by weight of dibutyl tin di-2-ethylhexoate WHAT WE CLAIM IS:— 0.5% of vinyltriacetoxysilane 0.1% of acetic acid to a garment which comprises 0.5% of tetrabutyltitanate (1) applying to a fabric comprising fibres the remainder being aliphatic hydrocarbon formed from a thermoplastic organic polysolvent. mer a curable organosiloxane in an amount The silicone pick-up was 1.6 per cent by of at least 0.5% by weight calculated on 105 weight. the weight of the fabric, The fabric was dried, sewn into trousers (2) forming the fabric into a garment, and and pressed as shown in Example 2 and then (3) arranging those areas of the garment heated for 10 minutes at 163°C. The fabric

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showed excellent crease retention after five

Example 4

1.29% of a dimethylpolysiloxane liquid hav-

ing a viscosity of 100,000 cs. at 25°C. 0.44% of methyl hydrogen polysiloxane

The fabric of Example 2 was padded with

washings as shown above.

an emulsion containing:

1. A process for imparting durable creases 100

in which it is desired to form creases in creases and subjecting the garment, to 110 sufficient heat and pressure whilst so arranged so as to set the thermoplastic fibres and cure the siloxane to a flexible resin, said siloxane being the only permanent crease-forming resin (as hereinbefore 115 defined) applied to the fabric.

2. A process as claimed in claim 1 in which the siloxane is a dimethylsiloxane.

3. A process as claimed in claim 1 substantially as described with reference to any of the Examples.

of the Examples.

4. A garment prepared by the process claimed in any of the preceding claims.

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